

In re Avandia Marketing, Sales Practices and Products..., Not Reported in...

2012 WL 3205620

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United States District Court,  
E.D. Pennsylvania.

In re AVANDIA MARKETING, SALES  
PRACTICES AND PRODUCTS LIABILITY  
LITIGATION.

This Document Applies to:  
Amjad Faheem v. GlaxoSmithKline, LLC  
Marvin Rainey v. GlaxoSmithKline, LLC

MDL No. 1871. | No. 07-MD-01871. | Civil Action  
Nos. 11-695, 11-3031. | Aug. 7, 2012.

**Opinion**

RUFE, District Judge.

\*<sup>1</sup> Plaintiffs in these cases filed suit alleging that they suffered heart-related injuries caused by their ingestion of the drug Avandia. Defendant, GlaxoSmithKline, LLC (“GSK”), has filed motions for summary judgment, contending that Plaintiffs’ claims are barred by the applicable statutes of limitations.<sup>1</sup> Plaintiffs, through the Plaintiffs’ Steering Committee (“PSC”), oppose the motions and have moved for additional discovery pursuant to Federal Rule of Civil Procedure 56(d).<sup>2</sup>

**I. BACKGROUND**

Plaintiff Marvin Rainey, a resident of Tennessee, began using Avandia in 1999 and suffered a heart attack in 2000; Plaintiff Amjad Faheem, a resident of Kentucky, began using Avandia in 2001 and suffered a heart attack in 2004. Both Plaintiffs filed suit in 2011, alleging that their use of Avandia caused their injuries. Avandia, the brand name for rosiglitazone maleate, was approved by the Food and Drug Administration in 1999 and is manufactured by Defendant GSK. Avandia is a member of a class of drugs known as thiazolidinediones (“TZDs”), used to manage non-insulin-dependent diabetes, or Type 2 diabetes.

Defendant GSK seeks summary judgment on the statute of limitations as to these two Plaintiffs, but also seeks significantly broader relief. Specifically, GSK seeks to establish a “bar date,” i.e., the date by which any plaintiffs may be presumed as a matter of law to have been on notice of a possible link between Avandia and their

injuries, and therefore to pursue any tort claims. GSK argues that for plaintiffs alleging heart-related injuries from use of Avandia, the bar date is November 14, 2007.

**II. STANDARD OF REVIEW**

Upon motion of a party, summary judgment is appropriate if “the materials in the record” show “that there is no genuine dispute as to any material fact and the movant is entitled to judgment as a matter of law.”<sup>3</sup> Summary judgment may be granted only if the moving party persuades the district court that “there exists no genuine issue of material fact that would permit a reasonable jury to find for the nonmoving party.”<sup>4</sup> A fact is “material” if it could affect the outcome of the suit, given the applicable substantive law.<sup>5</sup> A dispute about a material fact is “genuine” if the evidence presented “is such that a reasonable jury could return a verdict for the nonmoving party.”<sup>6</sup>

In evaluating a summary judgment motion, a court “must view the facts in the light most favorable to the non-moving party,” and make every reasonable inference in that party’s favor.<sup>7</sup> Further, a court may not weigh the evidence or make credibility determinations.<sup>8</sup> Nevertheless, the party opposing summary judgment must support each essential element of the opposition with concrete evidence in the record.<sup>9</sup> If the evidence is merely colorable, or is not significantly probative, summary judgment may be granted.<sup>10</sup> This requirement upholds the “underlying purpose of summary judgment [which] is to avoid a pointless trial in cases where it is unnecessary and would only cause delay and expense.”<sup>11</sup> Therefore, if, after making all reasonable inferences in favor of the non-moving party, the court determines that there is no genuine dispute as to any material fact, summary judgment is appropriate.<sup>12</sup>

\*<sup>2</sup> Plaintiffs have filed a motion for additional discovery pursuant to Rule 56(d), which is “the proper recourse of a party faced with a motion for summary judgment who believes that additional discovery is necessary before he can adequately respond to that motion.”<sup>13</sup> A properly filed motion must be accompanied by “a supporting affidavit detailing what particular information is sought; how, if uncovered, it would preclude summary judgment; and why it has not previously been obtained.”<sup>14</sup>

**III. DISCUSSION**

**A. Applicable Law**

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The rules of the Judicial Panel on MultiDistrict Litigation allow cases to be filed directly in this District and made part of the Avandia MDL, which Plaintiffs in these cases did.<sup>15</sup> The Court must determine whether to apply Pennsylvania law or the law of Plaintiffs' home states. The Court has concluded, as have other MDL courts, that such cases should be governed by the law of the states where Plaintiffs received treatment and prescriptions for Avandia.<sup>16</sup> This ruling will promote uniform treatment between those Plaintiffs whose cases were transferred into the MDL from their home states and those Plaintiffs who filed directly into the MDL. This holding is also consistent with Pennsylvania's choice-of-law rules, because "Pennsylvania applies a flexible rule which permits analysis of the policies and interests underlying the particular issue before the court and directs courts to apply the law of the state with the 'most interest in the problem.' "<sup>17</sup> In personal injury cases, that is the state where the injury occurred.<sup>18</sup>

Faheem's home state, Kentucky, employs a one year statute of limitations for personal injury cases.<sup>19</sup> Kentucky law also recognizes the "discovery rule," under which "[a] cause of action will not accrue ... until the plaintiff discovers or in the exercise of reasonable diligence should have discovered not only that he has been injured but also that his injury may have been caused by the defendant's conduct."<sup>20</sup> Reasonable diligence requires that the plaintiff be "as diligent as the great majority of persons would [be] in the same or similar circumstances...."<sup>21</sup>

Rainey's home state, Tennessee, also has a one-year statute of limitations in personal injury cases,<sup>22</sup> with the cause of action generally accruing on the date of the injury.<sup>23</sup> Tennessee also recognizes the discovery rule which tolls the statute of limitations until "one discovers, or in the exercise of reasonable diligence should have discovered, both (1) that he or she has been injured by wrongful or tortious conduct and (2) the identity of the person or persons whose wrongful conduct caused the injury."<sup>24</sup> This only requires that the plaintiff be aware of those facts sufficient "to place a reasonable person on notice that the injury was the result of the wrongful conduct of another."<sup>25</sup>

## **B. 2007 Evidence of a Possible Link between Avandia Use and Heart-Related Injuries**

### **1. The Nissen Study and FDA Action**

\*3 After conducting a meta-analysis study, Dr. Steven Nissen concluded that use of Avandia was associated with an increased risk of heart attack. Specifically, the Nissen study found that Avandia increased the risk of myocardial

infarction by 43%, a statistically significant result.<sup>26</sup> The New England Journal of Medicine published the peer-reviewed Nissen study on May 21, 2007. In response to the Nissen study's publication, the American College of Cardiology, American Diabetes Association, and American Heart Association issued a statement expressing concern and advising patients with diabetes to speak with their physicians.<sup>27</sup> At a meeting in July 2007, the Food and Drug Administration Advisory Committee voted 20–3 that "available data support a conclusion that Avandia increases cardiac ischemic risk," but did not act at that time to restrict the availability of Avandia.<sup>28</sup> However, the FDA did require that GSK revise the product label for Avandia, and GSK agreed to include within a black box warning the statement that "Avandia was not recommended for any patient with symptomatic heart failure," to add a summary of the results of an integrated data set from 42 clinical trials regarding risk of myocardial ischemic events, and to include more detailed results in the Warnings section in the label.<sup>29</sup>

### **2. "Dear Healthcare Professional" and "Dear Patient" Letters**

From May through November 2007, GSK sent a series of letters to healthcare professionals regarding studies of Avandia and cardiovascular health.<sup>30</sup> These letters discussed various studies, including the Nissen study (and GSK's disagreement with it)<sup>31</sup> as well as regulatory developments with regard to cardiovascular risk and Avandia use, culminating in a November 2007 letter reporting on the label revision. Any physician receiving these letters would be aware that there was concern about cardiovascular health and use of Avandia, although the letters expressed GSK's view that Avandia remained "an important treatment option for physicians" in treating diabetes.<sup>32</sup> On June 1, 2007, GSK also published a "Dear Avandia Patient" letter, which responded to the "recent press coverage about the safety of Avandia" and stated that GSK stood firmly behind Avandia.<sup>33</sup> Plaintiffs have produced evidence that during this same time frame, GSK criticized the Nissen study, and worked to encourage physicians to continue to prescribe Avandia.<sup>34</sup>

### **3. Media Reports**

The publication of the Nissen Study generated substantial interest in the media. Significantly, in the days following the publication, television news programs highlighted the findings of the Nissen report, in several instances as the lead story on the national evening broadcast.<sup>35</sup> These reports summarized the findings of the Nissen study and also noted that GSK "strongly disagrees with the

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conclusions ... and says other studies prove the drug's safety."<sup>36</sup> During the summer and fall of 2007, national and local newspapers published articles of varying depth and prominence discussing reported risks of Avandia use.<sup>37</sup> Many of these news reports included GSK's assurances that Avandia was safe and effective. The media also reported the actions of the FDA Advisory Committee, describing it in at least one report as sending a "mixed message" to Avandia patients.<sup>38</sup> The November 2007 label revision generated still more news stories, which included the information that the FDA had decided to keep Avandia on the market and noted that the evidence of an increased risk of cardiovascular events was "inconclusive."<sup>39</sup>

### **C. The Cumulative Effect of 2007 Events Triggered the Duty to Investigate**

\*4 The evidence shows that the events described above were regarded as significant by physicians, patients, and attorneys. By August 2007, Avandia prescriptions had fallen by 45%; by November 2007, sales had fallen 54%.<sup>40</sup> This MDL was formed in 2007 as a result of numerous lawsuits filed nationwide that year.

The question then becomes, did all of these events suffice, as a matter of law, to put on notice those who had suffered heart-related injuries that Avandia could be to blame and trigger a duty to investigate? The extensive media reports "indicate what was in the public realm at the time, not whether the contents of those articles were in fact true."<sup>41</sup> What was in the public realm throughout the second half of 2007 linked Avandia use with the possibility of heart-related illness, although the reports certainly did not reach an unqualified conclusion in that regard.<sup>42</sup>

The Court concludes that a reasonable person who knew that he or she had suffered cardiovascular injury and had taken Avandia would have been put on notice by the end of 2007 of the need to investigate a possible link between Avandia and the injury. Plaintiffs argue strenuously that GSK concealed information regarding the risks of Avandia and continued to downplay the seriousness of the risks until 2010, and that in August 2007, GSK argued to physicians that the "totality of evidence" showed "[n]o increased risk of [cardiovascular events] vs. oral antidiabetic agents."<sup>43</sup> Accepting Plaintiffs' argument as true for purposes of the motions for summary judgment, Plaintiffs in these cases *had already suffered heart attacks, and knew that they had done so*. A reasonable person who had suffered a heart attack, and who had taken Avandia, as Plaintiffs here did, would have been on notice by the end of 2007 to investigate a possible link

between the Avandia use and the heart attack.<sup>44</sup> Similarly, Plaintiffs' arguments of fraudulent concealment miss the mark. The issue of whether GSK should have disclosed more information or disclosed it sooner does not affect what information became available in 2007.

### **D. Plaintiffs' Rule 56(d) Motion**

Plaintiffs have moved for time to take additional discovery, including individualized discovery as to what Plaintiffs and their physicians knew and when, and extensive discovery that essentially relates to GSK's alleged fraudulent concealment. The Court finds that these are not typical cases where summary judgment is sought before discovery: during the course of the MDL, many hundreds of thousands of pages of documents have been produced, and Plaintiffs have not demonstrated that more is necessary on the issues discussed herein. Further, discovery as to the personal circumstances of Plaintiffs is not required because the evidence presented demonstrates as a matter of law that the information available both to the general public and to treating physicians throughout 2007 should have put a reasonable person on notice to investigate the possible link between a heart attack already suffered and use of Avandia.<sup>45</sup>

### **E. Limitations of the Court's Ruling**

\*5 The Court holds that under the laws of Tennessee and Kentucky, a reasonable person who knew that he or she had suffered a heart-related injury after taking Avandia was on notice by the end of 2007<sup>46</sup> to investigate the possibility of a link between Avandia and their injury so as to start the statute of limitations running on tort claims alleging personal injury. This ruling does not address Avandia patients who suffered other injuries, such as stroke; nor does it address any other claims asserted against GSK. The Court is not ruling at this time on whether GSK concealed evidence of the risks of Avandia use. The Court also notes that the law of certain states may have a different view of when a claim is tolled.

## **IV. CONCLUSION**

The Court holds that a reasonable person who knew that he or she had suffered a heart-related injury after taking Avandia was on notice to investigate the possible link between the injury and Avandia use by December 31, 2007. Because Plaintiffs did not file their personal-injury claims within the applicable statute of limitations, GSK's Motions for Summary Judgment will be granted, and those claims will be dismissed. An appropriate order will

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be entered.

Footnotes

- 1 GSK filed the Motion for Summary Judgment in five cases. By notice dated February 28, 2012, GSK withdrew the motion in the case of *Randall v. GlaxoSmithKline, LLC*, Civil Action No. 10-4861, as the case became subject to a pending settlement agreement. By letter dated June 6, 2012, GSK advised the Court that an agreement in principle had been reached to settle the cases of *Bonn v. GlaxoSmithKline, LLC*, Civil Action No. 11-2734, and *Estate of Henry v. GlaxoSmithKline*, Civil Action No. 10-4080.
- 2 By Memorandum and Orderdated September 7, 2011, the Court denied motions to dismiss based on the statute of limitations in 60 cases. Defendant GlaxoSmithKline, LLC (“GSK”) filed motions for partial reconsideration, seeking again to dismiss all or part of 49 of these cases; those motions were also denied. In the earlier rulings, the Court held that the Court could not make a determination in the context of a motion to dismiss, but required an evidentiary record.
- 3 Fed.R.Civ.P. 56(a), (c)(1)(A).
- 4 *Miller v. Ind. Hosp.*, 843 F.2d 139, 143 (3d Cir.1988).
- 5 See *Anderson v. Liberty Lobby, Inc.*, 477 U.S. 242, 248 (1986).
- 6 *Id.*
- 7 *Hugh v. Butler Cnty. Family YMCA*, 418 F.3d 265, 267 (3d Cir.2005).
- 8 *Boyle v. County of Allegheny*, 139 F.3d 386, 393 (3d Cir.1998).
- 9 *Celotex Corp. v. Catrett*, 477 U.S. 317, 322–23 (1986).
- 10 *Anderson*, 477 U.S. at 249–50 (citations omitted).
- 11 *Walden v. Saint Gobain Corp.*, 323 F.Supp.2d 637, 641 (E.D.Pa.2004) (citing *Goodman v. Mead Johnson & Co.*, 534 F.2d 566, 573 (3d Cir.1976)).
- 12 *Celotex*, 477 U.S. at 322; *Wisniewski v. Johns-Manville Corp.*, 812 F.2d 81, 83 (3d Cir.1987).
- 13 *Murphy v. Millennium Radio Grp. LLC*, 650 F.3d 295, 309 (3d Cir.2011).Rule 56(d) provides that “[i]f a nonmovant shows by affidavit or declaration that, for specified reasons, it cannot present facts essential to justify its opposition, the court may: (1) defer considering the motion or deny it; (2) allow time to obtain affidavits or declarations or to take discovery; or (3) issue any other appropriate order.”
- 14 *Doe v. Abington Friends Sch.*, 480 F.3d 252, 257 n. 3 (3d Cir.2007) (internal quotation omitted).
- 15 JPML Rule 7.2(a) provides that “[p]otential tag-along actions filed in the transferee district do not require Panel action. A party should request assignment of such actions to the Section 1407 transferee judge in accordance with applicable local rules.”
- 16 See, e.g., *In re Yasmin and Yaz (Drospirenone) Mktg., Sales Practices & Prods. Liab. Litig.*, No. 3:09-MD-2100, 2011 WL 1375011, \*5 (S.D.Ill. Apr. 12, 2011) (holding that cases that originated outside of the court’s judicial district and that were filed directly into the MDL would be treated as if they were transferred from a judicial district sitting in the state where the case originated).
- 17 *Specialty Surfaces Int’l v. Cont’l Cas. Co.*, 609 F.3d 223, 229 (3d Cir.2010).

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18 See, e.g., *Flamer v. New Jersey Transit Bus Operations, Inc.*, 607 A.2d 260, 264 (Pa.Super.Ct.1992) (internal citation omitted). The Court does note that Pennsylvania has a “borrowing statute,” which provides that “[t]he period of limitation applicable to a claim accruing outside this Commonwealth shall be either that provided or prescribed by the law of the place where the claim accrued or by the law of this Commonwealth, whichever first bars the claim.”<sup>42</sup> Pa. Cons.Stat. Ann. § 5521(b). However, because the Court considers that “direct filed” cases should be treated as if they were filed in the Plaintiffs’ home states, the forum-shopping concerns of this statute are not implicated here.

19 See Ky.Rev.Stat. § 413.140(1), (1)(a).

20 *R.T. Vanderbilt Co., Inc. v. Franklin*, 290 S.W.3d 654, 659 (Ky.Ct.App.2009) (citing *Louisville Trust Co. v. Johns-Manville Prods. Corp.*, 580 S.W.2d 497, 501 (Ky.1979) (quotation in *Louisville Trust* omitted)); see also *Johnson v. Sandoz Pharm. Corp.*, 24 F. App’x 533, 535–39 (6th Cir.2001) (applying Kentucky law in determining how the discovery rule affected the statute of limitations in products liability case where plaintiff claimed Parlodel led to stroke).

21 *Id.* (citing *Blanton v. Cooper Indus.*, 99 F.Supp.2d 797, 802 (E.D.Ky.2000) (quoting *Sawyer v. Midelfort*, 595 N.W.2d 423, 439 (Wis.1999)) (internal quotations omitted)).

22 See Tenn.Code Ann. § 28–3–104(a), (a)(1) (“Actions for ... injuries to the person ....“ “shall be commenced within one (1) year ....”).

23 Tenn.Code Ann. § 28–3–104(b)(1).

24 *Sherrill v. Souder*, 325 S.W.3d 584, 595 (Tenn.2010). See also *Wyatt v. A-Best, Co., Inc.*, 910 S.W.2d 851, 854 (Tenn.1995); *Potts v. Celotex Corp.*, 796 S.W.2d 678, 680–81 (Tenn.1990) (“[T]he statute [of limitations] is tolled only during the period when the plaintiff had no knowledge at all that the wrong had occurred and, as a reasonable person, was not put on inquiry.”); *Teeters v. Currey*, 518 S.W.2d 512, 512–17 (Tenn.1974) (first adopting the discovery rule); *Carter v. Danek Med. Inc.*, 1999 WL 33537317, at \*3–4 (W.D.Tenn.1999) (discussing the tolling of the statute of limitations in products liability claims involving spinal surgery).

25 *Id.*

26 *In re Avandia*, No. 07–1871, 2011 WL 13576, at \*3 (E.D.Pa. Jan. 4, 2011).

27 GSK Ex. 138.

28 PSC Ex. 124 at 4.

29 GSK Ex. 137.

30 GSK Exs. 130–37.

31 GSK Ex. 130 at 1.

32 E.g., GSK Ex. 137 at 2.

33 Compls. ¶ 55.

34 PSC Exs. 134–42.

35 GSK Exs. 9–19.

36 See, e.g., GSK Ex. 9.

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37 GSK Exs. 20–142. GSK also produced a listing of news reports that mentioned Avandia. GSK Exs. 5–6. The Court finds these exhibits of limited use to the Court; a number of the references appear to be to articles mentioning the effect of Avandia issues on GSK’s stock price, for example, and do not appear likely to have drawn general notice.

38 GSK Ex. 82 (ABC News transcript, July 31, 2007).

39 See, e.g., GSK Ex. 121 (New York Times article, Nov. 15, 2007).

40 PSC Ex. 128; GSK Rao Decl. Ex. 4.

41 *Benak ex rel. Alliance Premier Growth Fund v. Alliance Capital Management L.P.*, 435 F.3d 396, 401 n. 15 (3d Cir.2006).

42 Other Courts have set similar limitations periods. See, e.g ., *In re Briscoe*, 448 F.3d 201, 220–21 (3d Cir.2006) (in diet drugs litigation, finding that the statute of limitations barred claims after class notifications that followed withdrawal of the drugs from the market); *In re Vioxx Prods. Liab. Litig.*, 522 F.Supp.2d 799, 803 (E.D.La.2007) (finding that the statute of limitations barred claims after extensive publicity following the withdrawal of Vioxx from the market). Other courts have found that the statute of limitations applies even when a drug has not been withdrawn from the market. See, e.g., *In re Zyprexa Prods. Liab. Litig.*, 489 F.Supp.2d 230 (E.D.N.Y.2007).

43 PSC Ex. 126.

44 The Court notes that Type 2 diabetes is not a temporary condition; there is no cure, and patients who were receiving treatment in 2000 would still need to be managing their condition in 2007 (although not necessarily with medication).See Mayo Clinic Staff “Type 2 Diabetes” retrieved from <http://www.mayoclinic.com/health/type-2-diabetes/DS00585> (last viewed Aug. 1, 2012).

45 In addition, Plaintiffs were not prevented from offering evidence of their individual circumstances, for example, through affidavits.

46 Although GSK argues that the bar date should be November 14, 2007, the Court finds that news coverage of the 2007 label revision continued after that date, and therefore concludes that the last date on which Plaintiffs should have been on notice is December 31, 2007.

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**Sanchez v. Boston Scientific Corp., Slip Copy (2014)**

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United States District Court, S.D. West Virginia.

Roseanne SANCHEZ, et al., Plaintiffs,  
v.  
BOSTON SCIENTIFIC CORPORATION,  
Defendant.

Civil Action No. 2:12-cv-05762. | Jan. 17, 2014.

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**MEMORANDUM OPINION AND ORDER**

**(Boston Scientific's Motion for Summary Judgment  
Based on the Statute of Limitations)**

[JOSEPH R. GOODWIN](#), District Judge.

\*1 Pending before the court is Boston Scientific Corporation's Motion for Summary Judgment Based on the Statute of Limitations [Docket 30]. Relying on California's two-year statute of limitations, Boston Scientific argues that Ms. Sanchez's claim is time-barred. In its supporting memorandum, Boston Scientific states that Ms. Sanchez underwent four revision surgeries more than two years before she filed this action. Boston Scientific claims these surgeries put Ms. Sanchez on actual or inquiry notice of her claim more than two years before filing suit. For the reasons stated below, Boston

Boston Scientific's motion for summary judgment [Docket 30] is **DENIED**.

**I. Background**

This case is one of several thousand assigned to me by the Judicial Panel on Multidistrict Litigation and one of four (now three) bellwether cases set for trial pursuant to Pretrial Order # 54 [Docket 22]. These cases involve the use of transvaginal surgical mesh to treat pelvic organ prolapse ("POP") and stress urinary incontinence ("SUI").

On January 13, 2010, Dr. Kerri Wiltchik, M.D., implanted Ms. Sanchez with a Pinnacle Pelvic Floor Repair Kit and an Advantage Transvaginal Mid-Urethral Sling System. (*See* Boston Scientific Corp.'s Mem. in Supp. of its Mot. for Summ. J. based on the Statute of Limitations, Exhibit A [Docket 30-1], at 85-87; Exhibit B [Docket 30-2], at 3).<sup>1</sup> The implantation surgery took place at Marian Medical Center in Santa Maria, California. (*See* Exhibit A [Docket 30-1], at 85). The products were implanted to treat Ms. Sanchez's SUI, POP, and cystocele. (Exhibit C [Docket 30-3], at 4).

According to Ms. Sanchez's plaintiff fact sheet, she first saw a health care provider for symptoms related to the mesh in February 2010. (Exhibit C [Docket 30-3], at 6). In addition, Ms. Sanchez's deposition testimony indicated that she was experiencing a pink-tinged discharge every day since the implantation surgery. (Exhibit E [Docket 30-5], Deposition of Roseanne Sanchez, at 21:1-15). Between her implantation surgery and her first revision surgery, Ms. Sanchez complained of vaginal discharge, itching, and abdominal cramping. (Exhibit A [Docket 30-1], at 50, 52).

On April 9, 2010, approximately four months after the implantation surgery, Ms. Sanchez told Dr. Wiltchik she was experiencing "abnormal vag[inal] bleeding scant with a pink discharge which causes her to wear a daily panty liner" and also felt "something scratchy like a stitch in her vagina." (Exhibit A [Docket 30-1], at 46). Dr. Wiltchik diagnosed Ms. Sanchez as having "complications due to genitourinary device, graft, and implant." *Id.* Dr. Wiltchik excised a small portion of the mesh and applied silver nitrate to the area. (Exhibit D [Docket 30-4], Deposition of Dr. Kerri Wiltchik, at 50:1-3). Dr. Wiltchik prescribed Vagifem tablets, which would help grow the mucosa over the exposed mesh areas and promote healing. (*Id.* at 185-86:24-3). Ms. Sanchez's medical records for that day indicate she understood "that a few treatments may be required before the exposed mesh areas are completely

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covered and her symptoms resolve."(Exhibit A [Docket 30-1], at 46).

\*2 On May 3, 2010 Dr. Wiltchik performed a second revision surgery. (Exhibit A [Docket 30-1], at 44). Dr. Wiltchik again concluded that Ms. Sanchez was suffering from "complications due to genitourinary device, graft, and implant," specifically, exposed mesh from the Pinnacle product. (*Id.*; Exhibit D [Docket 30-4], Deposition of Dr. Kerri Wiltchik, at 49:19-22).

By her May 20, 2010, visit with Dr. Wiltchik, Ms. Sanchez testified that she was experiencing pelvic cramping and discomfort, which she believed were related to vaginal infections, as well as incontinence symptoms. (Exhibit E [Docket 30-5], Deposition of Roseanne Sanchez, at 225:11-15). Dr. Wiltchik again assessed that Ms. Sanchez's symptoms stemmed from complications with the pelvic implants. (Exhibit A [Docket 30-1], at 43). Dr. Wiltchik prescribed Metrogel-Vaginal gel, which Ms. Sanchez testified did not improve her symptoms. (*Id.* at 43; Exhibit E [Docket 30-5], Deposition of Roseanne Sanchez, at 224:14-17).

On June 14, 2010, Ms. Sanchez again complained to Dr. Wiltchik that she was experiencing copious amounts of pink-tinged discharge. (Exhibit A [Docket 30-1], at 41). According to Ms. Sanchez's medical records, "her discharge was thought to be due to her exposed mesh."(*Id.*). After a lengthy discussion with Dr. Wiltchik, Ms. Sanchez agreed to undergo another revision surgery, this time under general anesthesia. (*Id.* at 42). Ms. Sanchez understood that the procedure would help stop the mesh from poking through her vaginal wall. (Exhibit E [Docket 30-5], Deposition of Roseanne Sanchez, at 230:3-6). On June 18, 2010, Dr. Wiltchik removed a large portion of exposed mesh. (Exhibit A [Docket 30-1], at 80). This was Ms. Sanchez's third revision surgery.

Despite these three revisions of the mesh and other treatments, Ms. Sanchez's symptoms did not improve. (*See id.* at 39, 35). On September 1, 2010, Ms. Sanchez reported to Dr. Wiltchik that she was experiencing abnormal vaginal bleeding, pink-tinged discharge, and discomfort with intercourse. (*Id.* at 35). The medical record for this date indicates Ms. Sanchez understood that "her symptoms are due to a small amount of exposed mesh."(*Id.*). For the fourth time, Dr. Wiltchik completed an in-office excision of the exposed mesh. (*Id.*). Later, on September 17, 2010, Ms. Sanchez agreed to undergo another revision surgery under anesthesia because her symptoms had not resolved. (*Id.* at 33).

According to the plaintiffs, during these medical visits,

Dr. Wiltchik never told Ms. Sanchez that her symptoms were related to a defect in the mesh. (*See Pls.' Resp. in Opp'n to Boston Scientific Corp.'s Mot. for Summ. J. based on the Statute of Limitations [Docket 32], at 4*). Ms. Sanchez testified that during one of her medical appointments, Dr. Wiltchik said, "[f]or one reason or another ... the skin was not healing over the mesh."(Exhibit E [Docket 30-5], Deposition of Roseanne Sanchez, at 221:11-13). According to Dr. Wiltchik's progress notes on May 11, 2011, she "discussed at length patient's reaction to mesh and propensity for body to expel mesh."(*See Pls.' Resp. in Opp'n to Boston Scientific Corp.'s Mot. for Summ. J. based on the Statute of Limitations, Exhibit 3 [Docket 32-3], Deposition of Dr. Kerri Wiltchik, at 191:9-14*).<sup>2</sup> Dr. Wiltchik told Ms. Sanchez she had "no idea why this was happening and for some reason [Ms. Sanchez's] body did not like" the mesh products. (*Id.* at 191:19-21). In addition, Dr. Wiltchik testified that she has never attributed the cause of Ms. Sanchez's symptoms to a defect in the mesh. (*See id.* at 223:17-21; 224:14-16, 23-25; 225:1-11).

\*3 Ms. Sanchez's plaintiff fact sheet indicates she became aware that her injuries were related to a defect in the mesh implants in August 2011. (Exhibit C [Docket 30-3], at 6). According to the plaintiffs, Ms. Sanchez saw an advertisement for transvaginal mesh litigation on television, which caused her to seek representation. (*See Exhibit 7 [Docket 32-7], at 28:5-16*). However, Ms. Sanchez's deposition testimony reveals that she did not know the month or the year she saw the advertisement. (Exhibit A [Docket 34-1], at 36:10-16). On September 21, 2012, Ms. Sanchez directly filed suit in MDL 2326 pursuant to Pretrial Order # 12 [Docket 176].<sup>3</sup>

## **II. Choice of Law**

In multidistrict litigation cases, the choice-of-law determination for pre-trial motions hinges upon whether federal or state law governs. "When analyzing questions of federal law, the transferee court should apply the law of the circuit in which it is located. When considering questions of state law, however, the transferee court must apply the state law that would have applied to the individual cases had they not been transferred for consolidation ." *In re Temporomandibular Joint (TMJ) Implants Prods. Liab. Litig.*, 97 F.3d 1050, 1055 (8th Cir.1996) (internal citations omitted); *see Toll Bros., Inc. v. Dryvit Sys., Inc.*, 432 F.3d 564, 568 n.4 (4th Cir.2005) (applying Connecticut state law in transferred multidistrict litigation case based on diversity jurisdiction and citing to *In re Temporomandibular (TMJ) Joint Implants Prods. Liab. Litig.*, 97 F.3d at 1055); *Bradley v. United States*, 161 F.3d 777, 782 n.4 (4th Cir.1998); *see*

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*also* Charles Alan Wright & Arthur R. Miller, *Federal Practice and Procedure* § 3866 (3d ed.2009).

This case is based on diversity jurisdiction. Federal law thus controls procedural issues and state law controls substantive issues. *Dixon v. Edwards*, 290 F.3d 690, 710 (4th Cir.2002). The standard for summary judgment is procedural; therefore, the federal standard applies. *Gen. Accident Fire & Life Assurance Co. v. Akzona, Inc.*, 622 F.2d 90, 93 n.5 (4th Cir.1980). In determining which state substantive law governs this dispute, I must first identify which choice-of-law rules to follow.

A majority of cases in an MDL are transferred from other forums pursuant to 28 U.S.C. § 1407. See William B. Rubenstein, *Newberg on Class Actions* § 10:29 (5th ed.2013). With respect to these transferred cases, courts routinely apply the choice-of-law of the originating forum. See, e.g., *In re Temporomandibular (TMJ) Joint Implants Prods. Liab. Litig.*, 97 F.3d at 1055; see also *Chang v. Baxter Healthcare Corp.*, 599 F.3d 728, 732 (7th Cir.2010) (“When a diversity case is transferred by the multidistrict litigation panel, the law applied is that of the jurisdiction from which the case was transferred....”).

However, plaintiffs may bypass the transfer process by directly filing into the MDL. See Andrew D. Bradt, *The Shortest Distance: Direct Filing and Choice of Law in Multidistrict Litigation*, 88 Notre Dame L.Rev. 759, 794 (2012). Some cases are directly filed into the MDL and originate in the MDL court’s judicial district. Others cases originate elsewhere and are directly filed into the MDL. See *In re Yasmin & Yaz (Drospirenone) Mktg., Sales Practices & Prods. Liab. Litig.*, MDL No. 2100, 2011 WL 1375011, at \*4 (S.D.Ill. Apr. 12, 2011).

\*4 The difficulty with the latter category of cases is there technically is no prior proper forum whose choice-of-law rules should apply. In addition, many direct filing orders indicate direct filing does not make the MDL court a “transferor court,” and thus has no effect on choice-of-law. Bradt, *supra*, at 764; see, e.g., Pretrial Order # 14 [Docket 196], at 3 (“This court shall not be deemed to be the ‘transferor court’ simply by virtue of the action having been directly filed into MDL No. 2326.”). Without a prior proper forum and a disclaimer that direct filing does not affect choice-of-law, it may be difficult to determine which forum’s choice-of-law should apply.

For cases that originate outside the MDL court’s judicial district and are filed directly into the MDL, many courts apply the choice-of-law rules of the “originating jurisdiction.” *In re Watson Fentanyl Patch Prods. Liab. Litig.*, MDL No. 2732, 2013 WL 4564927, at \*2 (N.D.Ill.

Aug. 27, 2013) (“Indeed, the prevailing rule in this situation is that in a case that was directly filed in the MDL transferee court but that originated elsewhere, the law (including the choice of law rules) that applies is the law of the state where the case originated.”); *In re Yasmin & Yaz (Drospirenone) Mktg., Sales Practices & Prods. Liab. Litig.*, 2011 WL 1375011, at \*6 (“[T]he Court concludes that the better approach is to treat foreign direct filed cases as if they were transferred from a judicial district sitting in the state where the case originated.”). See generally *Wahl v. Gen. Elec. Co.*, No. 3:13-CV-0329, 2013 WL 604818, at \*4 (M.D.Tenn. Nov. 14, 2013) (“Most of the courts that have considered this peculiar procedural posture have stated that it is appropriate to apply the choice of law rules of the ‘originating’ jurisdiction (i.e., where the case would have [been] brought but for the CMO permitting direct filing), rather than the choice of law rules of the MDL Court.”).

In prescription drug MDLs, the originating jurisdiction is the place where the drug was purchased and prescribed. *In re Yasmin & Yaz (Drospirenone) Mktg., Sales Practices & Prods. Liab. Litig.*, 2011 WL 1375011, at \*6 (“[T]he better approach is to treat foreign direct filed cases as if they were transferred from a judicial district sitting in the state where the case originated,” which is “the state where the plaintiff purchased and was prescribed the subject drug.”); *In re Avandia Mktg., Sales Practices & Prods. Liab. Litig.*, MDL No. 1871, 2012 WL 3205620, at \*2 (E.D.Pa. Aug. 7, 2010) (“The Court has concluded, as have other MDL courts, that such cases should be governed by the law of the states where Plaintiffs received treatment and prescriptions for Avandia. This ruling will promote uniform treatment between those Plaintiffs whose cases were transferred into the MDL from their home states and those Plaintiffs who filed directly in the MDL.”).

For cases that originate elsewhere and are directly filed into the MDL, I will follow the better-reasoned authority that applies the choice-of-law rules of the originating jurisdiction, which in our case is the state in which the plaintiff was implanted with the product. Here, the plaintiff was implanted with the product in Santa Maria, California. Therefore, California choice-of-law rules will govern the selection of the statute of limitations.

\*5 California uses the governmental interest approach to analyze choice-of-law questions. *Wash. Mut. Bank v. Superior Court*, 15 P.3d 1071, 1080–81 (Cal.2001). Under this approach, the parties agree that the California statute of limitations would apply. Because the parties agree on this point, I will assume that it is not an issue. Therefore, I will apply the California statute of limitations

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to determine whether Ms. Sanchez's claim is time-barred.

### **III. Legal Standard**

To obtain summary judgment, the moving party must show that there is no genuine issue as to any material fact and that the moving party is entitled to judgment as a matter of law. Fed.R.Civ.P. 56(a). In considering a motion for summary judgment, the court will not "weigh the evidence and determine the truth of the matter." *Anderson v. Liberty Lobby, Inc.*, 477 U.S. 242, 249 (1986). Instead, the court will draw any permissible inference from the underlying facts in the light most favorable to the nonmoving party. *Matsushita Elec. Indus. Co. v. Zenith Radio Corp.*, 475 U.S. 574, 587–88 (1986).

Although the court will view all underlying facts and inferences in the light most favorable to the nonmoving party, the nonmoving party nonetheless must offer some "concrete evidence from which a reasonable juror could return a verdict in his [or her] favor." *Anderson*, 477 U.S. at 256. Summary judgment is appropriate when the nonmoving party has the burden of proof on an essential element of his or her case and does not make, after adequate time for discovery, a showing sufficient to establish that element. *Celotex Corp. v. Catrett*, 477 U.S. 317, 322–23 (1986). The nonmoving party must satisfy this burden of proof by offering more than a mere "scintilla of evidence" in support of his or her position. *Anderson*, 477 U.S. at 252. Likewise, conclusory allegations or unsupported speculation, without more, are insufficient to preclude the granting of a summary judgment motion. See *Felty v. Graves Humphreys Co.*, 818 F.2d 1126, 1128 (4th Cir.1987); *Ross v. Comm'n's Satellite Corp.*, 759 F.2d 355, 365 (4th Cir.1985), abrogated on other grounds, *Price Waterhouse v. Hopkins*, 490 U.S. 228 (1989).

### **IV. Discussion**

In California, there is a two-year statute of limitations for personal injury actions. Cal.Civ.Proc.Code § 335.1. This statute applies to injuries involving defective products regardless of the legal theory asserted. See, e.g., *Soliman v. Philip Morris Inc.*, 311 F.3d 966, 971 (9th Cir.2002). The general rule is that a cause of action accrues when all of its elements are complete. *Fox v. Ethicon Endo-Surgery, Inc.*, 110 P.3d 914, 920 (Cal.2005). However, the discovery rule tolls accrual until the plaintiff "is aware of her injury and its negligent cause." *Jolly v. Eli Lilly & Co.*, 44 Cal.3d 1103, 1109 (1988). In other words, the statute begins to run "when the plaintiff suspects or should suspect that her injury was

caused by wrongdoing, that someone has done something wrong to her." *Id.* at 1110. "Wrong" is not used "in any technical sense, but rather in accordance with a lay understanding." *Id.* "The question when a plaintiff actually discovered or reasonably should have discovered the facts for purposes of the delayed discovery rule is a question of fact unless the evidence can support only one reasonable conclusion." *Ovando v. County of Los Angeles*, 159 Cal.App. 4th 42, 61 (2008).

\*6 In a case involving both medical malpractice and products liability claims, the California Supreme Court has stated that "a plaintiff's ignorance of wrongdoing involving a product's defect will usually delay accrual because such wrongdoing is essential to that cause of action." *Fox*, 110 P.3d at 924. Simply put, "[t]he discovery rule does not trigger accrual of a cause of action unless the plaintiff has some reason to suspect wrongdoing; that is, when a plaintiff, through reasonably diligent investigation, discovers only that he has been injured but not that the injury may have a wrongful cause, then the clock has not yet begun to run." *Hendrix v. Novartis Pharm. Corp.*, No. CV-13-2402-MWF PLAX, 2013 WL 5491846, at \*5 (C.D.Cal. Oct. 2, 2013).

However, in order to use the discovery rule to delay accrual, the plaintiff must "plead facts to show (1) the time and manner of discovery and (2) the inability to have made earlier discovery despite reasonable diligence." *Fox*, 110 P.3d at 921. In other words, "to employ the discovery rule to delay accrual of a cause of action, a potential plaintiff who suspects that an injury has been wrongfully caused must conduct a reasonable investigation of all potential causes of that injury. If such an investigation would have disclosed a factual basis for a cause of action, the statute of limitations begins to run on that cause of action when the investigation would have brought such information to light." *Id.*

Boston Scientific argues that Ms. Sanchez had discovered the wrongful cause of her injuries more than two years before she filed suit. First, Ms. Sanchez had at least four revision surgeries, which should have put her on actual or inquiry notice that her symptoms were related to a problem with the mesh implants. In support of this contention, Boston Scientific cites *Coleman v. Boston Scientific*, No. 1:10-CV-01968-0WW, 2011 WL 3813173 (E.D.Cal. Aug. 29, 2011). In *Coleman*, the plaintiff was implanted with a Boston Scientific mesh product on December 5, 2006. *Id.* at \*1. "From July 2007 to March 2009, the plaintiff had surgery, vaginal reconstruction, and mesh removal in order to treat her 'recurrent symptoms of pelvic pain, erosion and recurrent infection of the tissue surrounding the mesh.' " *Id.* The

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plaintiff filed suit more than two years after her July 2007 surgery. *See id.* The plaintiff argued she was not on notice until she had seen a 2008 FDA Notice regarding the possible defectiveness of the product. *Id.* at \*3. The United States District Court for the Eastern Division of California stated that

A reasonable person who is implanted with a medical device, which requires a second corrective surgery to remove the device and correct injuries resulting there from within a year of implantation [,] should suspect the defectiveness of the device and conduct a reasonable inquiry and examination into the suitability of the device.

*Id.* Second, Boston Scientific claims that if Ms. Sanchez had consulted Dr. Wiltchik, her medical records, or the publicly available 2008 FDA Notice, she would have discovered that her injuries were related to a problem with the mesh.

\*7 The plaintiff counters that *Coleman* does not apply to this case because the court never found as a matter of law that the plaintiff had discovered the wrongful cause of her injury. In a footnote, the court noted that “[d]ue to the lack of detail concerning the nature of Plaintiff’s 2007 surgery, it cannot be said as a matter of law that Plaintiff was on inquiry notice of her claims in 2007.” *Id.* at \*3 n.1. Instead, the plaintiffs claim Ms. Sanchez had notice of the wrongful cause of her symptoms, i.e. product defect, when she observed a television advertisement about mesh litigation in August 2011. The plaintiffs argue that Ms. Sanchez did not initially suspect a defect caused her injuries because Dr. Wiltchik stated that her symptoms were related to her body’s rejection of the mesh, not the mesh itself. In addition, the plaintiffs contend that Ms. Sanchez’s nineteen visits with her physician, or an investigation of her medical records, would not have revealed this wrongful cause because Dr. Wiltchik did not suspect that Ms. Sanchez’s injuries were caused by a defect in the mesh.

In other words, the plaintiffs claim that summary judgment is not appropriate because the record supports two inferences: First, Ms. Sanchez initially suspected that her symptoms were related to her body’s rejection of the mesh and only in August 2011 did she discover her symptoms had a wrongful cause. Conversely, she knew after four surgeries, including a surgery under anesthesia, that something was wrong with the product, not her body, and thus was on inquiry notice.

For instruction on this point, the plaintiffs cite *Clark v. Baxter Healthcare Corp.*, 83 Cal.App. 4th 1048 (2000). In *Clark*, the plaintiff began experiencing allergic reactions to latex gloves starting in 1992. *Id.* at 1053. The plaintiff consulted with an allergist who suggested that the plaintiff was allergic to the gloves. *Id.* In May 1995, the plaintiff had a serious allergic response to the latex gloves. *Id.* at 1053. By the end of 1995, the plaintiff joined a support group. *Id.* The group gave her a flyer regarding latex allergies litigation, which indicated that there might be a defect in the latex gloves. *Id.* The plaintiff filed suit against the glove manufacturers on January 23, 1996. *Id.* The flyer was entered into the record, and the plaintiff submitted a declaration stating when she had received the flyer. *Id.* The defendants moved for summary judgment based on the statute of limitations. *Id.*

The California Court of Appeals found that there was a triable issue of fact because the record supported two reasonable inferences—(1) the plaintiff “could reasonably have inferred from the advice given her by various doctors and from the severity of the May 1995 acute reaction, caused by gloves she was not wearing, that more than a natural allergy to a natural substance was involved, and that a product defect or a contaminated product could have been a causative factor” or (2) “that she did not become aware of a potential wrongfulness component of her cause of action until more information than the existence of her allergies placed her on inquiry notice and then was actually gained.” *Id.* at 1059–60. If the plaintiff identified the negligent cause of her injuries by May 1995, her action would be time-barred. However, if she was unaware of this negligent cause until the end of 1995, her claim was timely filed. Accordingly, the court denied the defendants’ motion for summary judgment.

\*8 If I were permitted to weigh evidence at the summary judgment stage, it is unlikely that Ms. Sanchez would prevail. However, I am not a fact finder. Therefore, based on the record before me, I must reluctantly conclude that there is a genuine issue of material fact regarding when Ms. Sanchez suspected that wrongdoing caused her injuries.

On one hand, a jury might believe that Ms. Sanchez initially suspected the cause of her symptoms was related to her body’s rejection of the mesh. Dr. Wiltchik never told Ms. Sanchez that her symptoms were caused by a defect in the mesh. In addition, Dr. Wiltchik never suspected that a defect could be causing Ms. Sanchez’s symptoms. One could reasonably conclude that Ms. Sanchez’s body simply “didn’t like” the mesh. Ms. Sanchez may have continued to believe this to be the

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cause of her symptoms until August 2011, when she allegedly viewed the television advertisement. The jury might reach this conclusion even though Ms. Sanchez's deposition testimony reveals she could not remember the exact date when she saw the advertisement. Thus, a jury could reasonably infer that Ms. Sanchez discovered the wrongful cause of injuries in August 2011 and thus timely filed her action.

On the other hand, a jury might conclude that after four revision surgeries, several medical treatments, and nineteen medical appointments, her body's rejection of the mesh was not a reasonable explanation of her symptoms. A reasonable person could conclude that the cause of Ms. Sanchez's injuries was a defect in the product, not her body's natural reaction to the mesh. Thus, a jury could infer that Ms. Sanchez discovered the wrongful cause of her injuries more than two years before filing suit. Therefore, her claim could be time-barred.

Assuming that Ms. Sanchez did suspect or should have suspected wrongdoing more than two years before filing, and thus had a duty to investigate, whether that investigation was reasonable is a more difficult issue. *See Fox*, 110 P.3d at 921 ("[A] potential plaintiff who suspects that an injury has been wrongfully caused must conduct a reasonable investigation of all potential causes of that injury."); *Jolly*, 44 Cal.3d at 1111 ("Once the plaintiff has a suspicion of wrongdoing, and therefore an incentive to sue, she must decide whether to file suit or sit on her rights. So long as a suspicion exists, it is clear that the plaintiff must go find the facts; she cannot wait for the facts to find her."); *Nelson v. Indevus Pharm., Inc.*, 142 Cal.App. 4th 1202, 1206 (2006) ("When the cases are read in whole, rather than in isolated quotes, it is clear that a plaintiff's duty to investigate does not begin until the plaintiff actually has a reason to investigate. A plaintiff has reason to discover a cause of action when he or she has reason at least to suspect a factual basis for its elements. We look to whether the plaintiffs have reason to

at least suspect that a type of wrongdoing has injured them."(internal citations, quotations, and alterations omitted)).

\*9 Because Dr. Wiltchik testified that she never told Ms. Sanchez her symptoms were due to a defect in the mesh, a jury could reasonably conclude that consultations with Dr. Wiltchik or a review of the medical records would not have given Ms. Sanchez a reason for suspicion of wrongdoing. (Exhibit 3 [Docket 32-3], Deposition of Dr. Kerri Wiltchik, at 223:17-21; 224:14-16; 224-225:23-11). While the evidence presented at trial may ultimately lead to a finding by the jury that Ms. Sanchez had a duty to investigate based on her multiple surgeries, there is enough of a material dispute to render summary judgment inappropriate.

For these reasons, I cannot determine as a matter of law that Ms. Sanchez discovered her cause of action more than two years before filing suit. Accordingly, I **DENY** Boston Scientific's motion for summary judgment. *See Ward v. Westinghouse Canada, Inc.*, 32 F.3d 1405, 1408 (9th Cir.1994) (applying California law) (factual issue regarding when plaintiff suspected wrongdoing not suitable for summary judgment).*See generally Sylve v. Riley*, 15 Cal.App. 4th 23, 26 (1993) ("Whether reasonable diligence was exercised is generally a question of fact precluding summary judgment.").

## V. Conclusion

For the reasons discussed above, Boston Scientific's Motion for Summary Judgment [Docket 30] is **DENIED**.The court **DIRECTS** the Clerk to send a copy of this Order to counsel of record and any unrepresented party.

## Footnotes

- <sup>1</sup> Exhibits relating to Boston Scientific's memorandum in support of its motion for summary judgment shall be identified alphabetically.
- <sup>2</sup> Exhibits relating to the plaintiffs' response to Boston Scientific's motion for summary judgment will be referred to numerically.
- <sup>3</sup> Pretrial Order # 12 was amended by Pretrial Order # 14 [Docket 196] on September 26, 2012. Pretrial Order # 14 did not modify sections regarding direct filing into the MDL.

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United States District Court,  
S.D. Illinois.

In re YASMIN AND YAZ (DROSPIRENONE)  
MARKETING, SALES PRACTICES AND  
PRODUCTS LIABILITY LITIGATION.  
This Document Relates To: All Cases.

No. 3:09-md-02100-DRH-PMF. | MDL No. 2100.  
| April 12, 2011.

## I. INTRODUCTION

DAVID R. HERNDON, Chief Judge.

\*1 This matter is before the Court for the purpose of resolving choice of law considerations pertaining to the issues of attorney-client privilege and the work-product doctrine. To date, Bayer has produced just under 3 million documents (approximately 65 million pages), and has withheld 12,857 unique documents (18,808 documents counting duplicates) as shielded by attorney-client privilege. Of those unique documents, the Bayer defendants assert that 6,282 are also protected under the work-product doctrine. The plaintiffs have currently asserted a representative challenge with regard to 330 of the allegedly privileged documents. The Bayer defendants report that this challenge affects 243 unique documents, 52 of which are also allegedly protected under the work-product doctrine.

The parties have attempted to independently resolve the dispute over the 330 challenged documents but are at an impasse with regard to the standard that should be applied when assessing whether the documents are protected under either the attorney-client privilege or the work-product doctrine. The first step in resolving this matter includes assessing both vertical and horizontal choice of law questions. The parties have asked the Court to resolve these choice of law questions and to provide direction with respect to procedure for addressing the remainder of the disputed issues.

## II. BACKGROUND

This multidistrict litigation ("MDL") concerns the prescription drugs Yaz, Yasmin, and Ocella and involves cases originating in nearly every state in the country as well as the District of Columbia and the territory of Puerto Rico. The basis for the Court's jurisdiction in these coordinated proceedings is diversity. The claims brought by the plaintiffs are state based claims for which state law supplies the rule of decision. The Bayer defendants, however, have asserted several federally based defenses. As of the Last status conference, the Bayer defendants report that 5,998 MDL cases have been served.

The MDL cases originate from three sources: (1) cases filed directly in this MDL that originated in this Court's judicial district ("local cases"); (2) cases that have been transferred from another district court pursuant to 28 U.S.C. § 1407 ("transfer cases"); and (3) cases that originated outside of this Court's judicial district and that were filed directly into the MDL ("foreign direct filed cases") pursuant to the Court's direct filing Order (MDL 2100 Docs 669 (original direct filing order), 1137(amended direct filing order), & 1462 (second amended direct filing order)).

## III. ANALYSIS

### A. Work-Product Doctrine

The Bayer defendants contend that a portion of the documents are protected from discovery by the work-product doctrine. The work-product doctrine, established in *Hickman v. Taylor*, 329 U.S. 495, 67 S.Ct. 385, 91 L.Ed. 451 (1947), is now codified in the Fed.R.Civ.P. 26(b)(3): "[A] party may not discover documents and tangible things that are prepared in anticipation of litigation or for trial by or for another party or its representative." The work-product doctrine is governed by federal law—even where the basis of federal jurisdiction is diversity. See e.g., *Pyramid Controls, Inc. v. Siemens Indus. Automations, Inc.*, 176 F.R.D. 269, 276 (N.D.Ill.1997) (Alesia, J.).

\*2 Accordingly, with regard to documents allegedly protected under the work-product doctrine, federal law controls.

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**B. Attorney–Client Privilege****1. Relevant Authority****a. Federal Rule of Evidence 501**

Federal Rule of Evidence 501 provides the rule for determining which body of law governs matters of privilege. Thus, the Court's analysis begins with the text of Rule 501, which provides:

Except as otherwise required by the Constitution of the United States or provided by Act of Congress or in rules prescribed by the Supreme Court pursuant to statutory authority, the privilege of a witness, person, government, State, or political subdivision thereof shall be governed by the principles of the common law as they may be interpreted by the courts of the United States in the light of reason and experience. However, in civil actions and proceedings, with respect to an element of a claim or defense as to which State law supplies the rule of decision, the privilege of a witness, person, government, State, or political subdivision thereof shall be determined in accordance with State law.

Fed.R.Evid. 501.

**b. Rule 501: Legislative History, State Law Proviso, and Reliance on1 Advisory Committee Notes****i. Legislative History and the State Law Proviso**

Originally, Article V of the Preliminary Draft of the Evidence Rules as proposed by the Advisory Committee consisted of thirteen rules recognizing specific privileges. 23 Charles Alan Wright & Kenneth Graham, Jr., *Federal Practice And Procedure: Evidence* § 5421. The proposed rule also made no provision for the application of state privilege law in diversity actions and clearly indicated that state privilege law should be disregarded. *Id.* This attempt to federalize the law of privilege was highly controversial and Congress consequently became actively involved in the rulemaking process. See *Id.* Congress "completely rejected" the Advisory Committee's privilege scheme, *id.*, and adopted the current version of Rule 501. *Id.*

The current version of Rule 501 was drafted by the House Judiciary Committee and includes a state law proviso designed to require the application of state privilege law in proceedings governed by *Erie R. Co. v. Tompkins*, 304 U.S. 64, 58 S.Ct. 817, 82 L.Ed. 1188 (1938). 23 Charles Alan Wright & Kenneth Graham, Jr., *Federal Practice And Procedure: Evidence* § 5421; Fed.R.Evid. 501 Advisory Committee Notes, H.R.Rep. No.93–650. In rendering its decision, the Court is mindful of the rationale underlying Rule 501's state law proviso:

[The] rationale underlying the proviso is that federal law should not supersede that of the States in substantive areas such as privilege absent a compelling reason. The Committee believes that in civil cases in the federal courts where an element of a claim or defense is not grounded upon a federal question, there is no federal interest strong enough to justify departure from State policy.

\*3 Fed.R.Evid. 501, Advisory Committee Notes, H.R.Rep. No. 93–650.

The Court also finds the review of the congressional history of Rule 501, provided by the Third Circuit Court of Appeals in an opinion authored by then Chief Judge Seitz in *Samuelson v. Susen*, 576 F.2d 546 (3rd Cir.1978) to be instructive on the matter. In *Samuelson*, the Third Circuit Court of Appeals explained that the congressional intent in enacting Rule 501 was to effectuate "state substantive rights, laws and policies in controversies where there is no substantial federal interest" and to preserve "the domain of state privilege law." *Id.* at 550. The court also noted that the decision to apply state privilege law in diversity actions was supported by the following contentions:

- (1) privilege rules are and should continue to be considered substantive for Erie purposes;
- (2) privilege rules are outcome determinative;
- (3) where state law supplies the rule of decision, state rules of privilege should be applied because there is no federal interest substantial enough to justify departure from state policy; and
- (4) state policy regarding privilege should not be thwarted merely because of diversity jurisdiction, a situation which, if allowed, would encourage forum

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shopping.

*Id.* citing H.R. Rep. No. 650, 93rd Cong., 1st Sess. 9 (1973).

### **ii. Reliance on the Advisory Committee Notes**

The Advisory Committee Notes that accompany Rule 501 consist of three reports: (1) House Report Number 93–650; (2) Senate Report Number 93–1277; and (3) Conference Report Number 93–1597. House Report Number 93–650 was drafted by the House Judiciary Committee and accompanied the House’s version of Rule 501 when it was first sent to the House floor for consideration. 23 Charles Alan Wright & Kenneth Graham, Jr., *Federal Practice And Procedure: Evidence* § 5421. After approval by the House, the Senate Judiciary Committee considered the House’s version of Rule 501. *Id.* The Senate Judiciary Committee, in Senate Report Number 93–1277, criticized certain provisions of the House’s version of Rule 501 and proposed an amendment to the state law proviso contained therein. *Id.* The proposed amendment was approved in the Senate without any debate. *Id.*

The Senate’s version of Rule 501 was then considered by the Conference Committee. *Id.* The Conference Committee ultimately rejected the Senate’s version of Rule 501 opting instead to adopt Rule 501 as originally proposed by the House. *Id.* The Conference Committee’s commentary explaining its reasoning and addressing some of the concerns that had been expressed in Senate Report Number 93–1277 is contained in Conference Report Number 93–1597. The House’s version of Rule 501 was then adopted without further debate. *Id.*

Any reliance this Court places on the Advisory Committee Notes that accompany Rule 501 is done in light of the legislative history discussed above. Of particular note, is the fact that the commentary in Senate Report Number 93–1277 was made in reference to the Senate’s version of Rule 501 which was rejected by the Conference Committee and therefore may not be an accurate indicator of Congressional intent with regard to Rule 501 as it was ultimately enacted. See 23 Charles Alan Wright & Kenneth Graham, Jr., *Federal Practice And Procedure: Evidence* § 5434 n. 15 & n. 17 (criticizing courts for relying on the Senate Report as evidence of congressional intent).

### **c. Rule 501 and Diversity Cases**

\*4 Pursuant to Rule 501, “with respect to an element of a

claim or defense as to which State law supplies the rule of decision,” matters of privilege are to “be determined in accordance with State law.” Fed.R.Evid. 501. This proviso has consistently been interpreted as requiring application of state privilege law in diversity based actions. See 2 Paul R. Rice & John B. Corr, *Attorney–Client Privilege in the U.S.* § 12:16 (2010). See also *Dunn v. Washington County Hosp.*, 429 F.3d 689, 693 (7th Cir.2005) (in a federal question case the Appellate Court noted that Illinois “peer review” privilege would not apply because state privilege only applies in diversity based actions); Fed.R.Evid. 501, Advisory Committee Notes (“The proviso is designed to require the application of State privilege law in civil actions and proceedings governed by *Erie R. Co. v. Tompkins*, 304 U.S. 64 (1938)”).

When Rule 501 requires application of state privilege law and there are factual connections to more than one state, federal courts apply state choice of law rules to determine which state’s privilege law controls.<sup>1</sup> Generally, a federal court sitting in diversity applies the choice of law rules emanating from the state in which it sits. *Tanner v. Jupiter Realty Corp.*, 433 F.3d 913, 915 (7th Cir.2006). When a case is transferred, however, the transferee court applies the choice of law rules of the state in which the transferor court sits. *Chang v. Baxter Healthcare Corp.*, 599 F.3d 728, 732 (7th Cir.2010). There is no controlling authority addressing this matter with regard to cases that (1) are directly filed in an MDL pursuant to a direct filing order and (2) originated outside of the MDL court’s judicial district.<sup>2</sup>

### **d. Rule 501 and Cases Involving Federal and State Claims or Defenses**

#### **i. Federal question cases and pendent state law claims**

The Seventh Circuit has held that in federal question cases federal privilege law applies even when the information sought would also be relevant to a pendent state claim. *Memorial Hospital for McHenry County v. Shadur*, 664 F.2d 1058, 1061 n. 3 (7th Cir.1981). In so holding, the Appellate Court reasoned that “it would be meaningless to hold the communication privileged for one set of claims and not the other.” *Id.*<sup>3</sup>

#### **ii. Diversity cases with state based claims and one or more federally based defenses**

The Court’s research reveals no Seventh Circuit authority addressing the controlling privilege law in diversity actions based on state causes of action that also involve federally based defenses.

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The Advisory Committee Notes to Rule 501, however, provide some guidance in this matter. Pursuant to the Conference Committee's Report, "If an item of proof tends to support or defeat a claim or defense, or an element of a claim or defense, and if state law supplies the rule of decision for that claim or defense, then state privilege law applies to that item of proof." Fed.R.Evid. 501, Advisory Committee Notes, Conf. Rep. No. 93-1597. Conversely, in diversity cases "where a claim or defense is based upon federal law....federal privilege law will apply to evidence relevant to the federal claim or defense." *Id.*<sup>4</sup>

\*5 There is no clear congressional intent with regard to whether federal or state privilege law should govern when, in a diversity action, an item of proof is relevant to both federal and state elements. See 23 Charles Alan Wright & Kenneth Graham, Jr., *Federal Practice And Procedure: Evidence* § 5434 (although the Senate Judiciary Report proposes application of the law favoring reception of the evidence, the Conference Committee Report is silent on the issue).

## 2. Analysis

### a. Illinois choice of law principles do not control simply because this MDL Court is sitting in diversity in Illinois

Plaintiffs contend that, because this Court is sitting in diversity in Illinois, Illinois is the forum and Illinois choice of law principles control. Plaintiffs' argument is incorrect because this MDL involves thousands of cases, many of which did not originate in this judicial district. As noted above (section II), cases in this MDL originate from three sources: (1) cases filed directly in this MDL that originated in this Court's judicial district ("local cases"); (2) cases that have been transferred from another district court pursuant to 28 U.S.C. § 1407 ("transfer cases"); and (3) cases that originated outside of this Court's judicial district and that were filed directly into the MDL ("foreign direct filed cases") pursuant to the Court's direct filing Order (MDL 2100 Docs 669 (original direct filing order), 1137 (amended direct filing order), & 1462 (second amended direct filing order)).

In the event that state law governs privilege matters in this MDL, the governing choice of law rules will depend on each case's source of origin. As to the local cases, Illinois choice of law rules control. *Tanner v. Jupiter Realty Corp.*, 433 F.3d 913, 915 (7th Cir.2006). As to transfer cases, however, Illinois choice of law rules are not controlling.<sup>5</sup> Instead, the choice of law rules that govern

are the rules of the jurisdiction from which each case was transferred. *Chang v. Baxter Healthcare Corp.*, 599 F.3d 728, 732 (7th Cir.2010).

As to the foreign direct filed cases, the choice of law decision is not as clear. Foreign direct filed cases are filed in this Court pursuant to a direct filing order (Case Management Order Number 9).<sup>6</sup> This agreed order authorizes the direct filing of cases that originated outside of this judicial district. The order expressly provides that the parties' direct filing agreement will not impact the choice of law that otherwise would apply to the direct filed actions.

In general, direct filing orders are beneficial to both parties because they streamline the litigation and help to eliminate the judicial inefficiency involved in the MDL transfer process. See Eldon Fallon, *Bellweather Trials in Multidistrict Litigation*, 82 Tul. L.Rev. 2323, 2353 (2008) (the direct filing procedure "eliminates the judicial inefficiency that results from two separate clerk's offices having to docket and maintain the same case and three separate courts (the transferor court, the MDL Panel, and the transferee court) having to preside over the same matter."). However, direct filing orders also present difficult choice of law issues. See e.g., *In re Express Scripts, Inc., PBM Litig.*, MDL No. 1672, 2007 WL 4333380, at \* 1-2 (E.D.Mo. Dec. 7, 2007) (not reported) (Limbaugh, J.) (applying the choice-of-law rules of the MDL forum to foreign case that was directly filed in the MDL); *In re Bausch & Lomb Inc. Contacts Lens Solution Prods. Liab. Litig.*, MDL No. 1785, 2007 WL 3046682, at \*3 (D.S.C. Oct. 11, 2007) (Norton, J.) (noting that "it would be an odd result to subject plaintiffs to [the law of the MDL forum] simply because they took advantage of the direct filing procedure—a procedure that provides benefits to all parties and preserves judicial resources").

\*6 The Court concludes that the better approach is to treat foreign direct filed cases as if they were transferred from a judicial district sitting in the state where the case originated. For purposes of this analysis, the Court considers the originating state to be the state where the plaintiff purchased and was prescribed the subject drug. Thus, for a foreign direct filed member action involving a plaintiff that purchased and was prescribed the subject drug in Tennessee, the Court will treat that plaintiff's claims as if they were transferred to this MDL from a district court in Tennessee.

### b. Wholesale Application of Federal Attorney-Client Privilege Law

In the instant case, application of state attorney-client

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privilege is a daunting task that involves considering the choice of law principles of all fifty states (as well as the District of Columbia and Puerto Rico) (*see sections III(B)(1)(c) & III(B) (2)(a)*). The Bayer defendants argue that engaging in such an analysis would be overly burdensome and contend that an appropriate alternative is the wholesale application of federal law. In support of their argument, the Bayer defendants point to the procedure utilized in the *Vioxx* MDL and outlined in *In re Vioxx Products Liability Litigation*, 501 F.Supp.2d 789 (E.D. La 2007) (Fallon, J.).<sup>7</sup>

In the *Vioxx* litigation, a discovery dispute arose regarding the defendant's claim of attorney-client privilege over approximately 500,000 pages of documents. The *Vioxx* court appointed Professor Paul R. Rice, an expert in privilege law, to review a representative sample of 2,000 documents withheld by the defendant and to make recommendations on whether the defendant's claims of privilege should be upheld. *Id.* at 789–795. During this process, Professor Rice applied federal privilege law standards, taking into account (1) "the law of attorney-client privilege both in general and in the context of [the *Vioxx*] multidistrict litigation" and (2) standards "common to all definitions of the attorney-client privilege." *Id.* at 794–795.

In the instant case, the Court is reluctant to follow the path taken by the *Vioxx* MDL. The *Vioxx* MDL's jurisdiction was based on diversity and the claims asserted by the plaintiffs turned on questions of state law. The *Vioxx* decision, however, does not address its reasoning for applying federal privilege law or Rule 501's state law proviso.

As discussed above (*see sections III(B)(1)(b)(i); III(B)(1)(c); and III(B)(1)(d)*), Rule 501 is designed to require the application of state privilege law in actions based on questions of state law between citizens of different states. Nothing in Rule 501 indicates that federal privilege standards may be applied wholesale to state question diversity proceedings simply because application of state privilege standards would be overly burdensome. In fact, the rationale underlying Rule 501 and its congressional history, which emphasize the view that privilege law is substantive and that federal law should not supersede that of the state law in diversity actions absent a compelling reason,<sup>8</sup> militate against the wholesale application of federal law for purposes of convenience (*see section III(B)(1)(b)*).

\*7 Further, although the Seventh Circuit has held that federal law applies in federal question cases (even where the same evidence is relevant to a pendent state claim),

*Shadur*, 664 F.2d at 1061 n. 3, the Seventh Circuit has not had occasion to address the wholesale application of federal law in diversity actions premised on state law claims.

Finally, the Conference Committee Report that accompanies Rule 501 indicates that in diversity actions based on questions of state law, a bifurcated approach is contemplated; privilege matters relevant to state claims or defenses will be governed by state privilege law and privilege matters relevant to federal claims or defenses will be governed by federal privilege law (*see sections III(B)(1)(b)(i) & III(B)(1)(d)(ii)*).

Accordingly, for the reasons discussed above, the Court finds that: (1) privilege matters that are relevant to an element of a federal defense will be governed by federal privilege law; and (2) privilege matters that are relevant to an element of a claim or defense for which state law supplies the rule of decision will be governed by state privilege law.

### **c. Privileged Material Relevant to Both Federal and State Elements**

This Court's decision may result in a scenario where the same privileged material is deemed relevant both to a state claim and a federal defense. Rule 501 does not expressly address what law must be applied in such a scenario. The Senate Report proposes that in diversity actions when the same evidence is relevant to state and federal elements, the "rule favoring reception of the evidence should be applied." Fed.R.Evid. 501 Advisory Committee Notes S.R. No. 93–1277. The Conference Committee Report, however, is silent on the issue and Congressional intent is therefore unclear. Fed.R.Evid. 501 Advisory Committee Notes C.R. No. 93–1597.

The Seventh Circuit has not directly addressed the issue. However, in *Estate of Suskovich v. Anthem Health Plans of Virginia, Inc.*, 553 F.3d 559, 570 (7th Cir.2009), the Appellate Court indicated that it would favor application of the approach proposed by the Senate Judiciary Committee. See *Estate of Suskovich*, 553 F.3d at 570 (noting with approval the district court practice of applying Federal Rule of Evidence 601, which creates a broad presumption of competency, when the evidence relates to state and federal claims and finding that the practice is consistent with the approach proposed by the Senate Judiciary Committee in Rule 501's Advisory Committee Notes). In addition, as discussed in section III(B)(1)(d)(i) above, in *Memorial Hospital for McHenry County v. Shadur*, 664 F.2d 1058, 1061 n. 3 (7th Cir.1981), the Appellate Court held that in a federal

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question case, where the same evidence is relevant to both state and federal elements, federal privilege law should govern because “it would be meaningless to hold the communication privileged for one set of claims and not the other.”*Id.*

\*8 Considering the above, the Court concludes that if the allegedly privileged material is relevant to both federal and state elements, the privilege law favoring admission shall govern (assuming of course there is a difference between the relevant federal and state privilege laws).

**d. When state law applies, section 139 of the Second Restatement will govern choice of law considerations**

As noted, to determine which state’s privilege law governs, the Court must look to state choice of law principles. In the instant case, this requires considering the choice of law principles applicable in all fifty states, the District of Columbia, and Puerto Rico. After conducting this analysis<sup>9</sup> and for the reasons outlined below, the Court concludes that section 139 of the Restatement (Second) Conflict of Laws (1971) (“Second Restatement”) will apply to privilege matters governed by state law in this MDL.

The Court’s research reveals that a majority of states have not established a choice of law doctrine regarding privileges. Courts that have analyzed conflicting privilege laws, however, tend to favor application of the most significant relationship test found in section 139 of the Second Restatement or of a similar test favoring application of the law of the state with the most significant relationship with the privileged communication—such as the law of the state where the communication is centered.

Of those jurisdictions that do not appear to have considered the issue (and therefore have not had occasion to accept or reject section 139), the majority (1) apply the Second Restatement to one or more other areas of law or (2) apply an interest based analysis that reflects the principles underlying section 139 of the Second Restatement. Given that these jurisdictions do not have a firmly established rule with regard to resolving privilege choice of law matters and these jurisdictions have applied the Second Restatement or a similar interest based analysis to other conflict of law issues, the Court concludes that, if presented with the issue, courts in these jurisdictions would apply the privilege law of the state with the most significant relationship to the communication and would be guided in that decision by the principles found in section 139 of the Second Restatement.

Finally, a small minority of jurisdictions continue to follow the traditional choice of law principles largely reflected in the original Restatement (First) of Conflict of Laws (1934) (“First Restatement”). The First Restatement, however, does not specifically address how privilege choice of law matters should be resolved. Under the First Restatement, matters are either substantive or procedural and procedural matters are to be governed by the law of the forum. Sections 596 and 597 of the First Restatement, specifically provide that the law of the forum governs issues of “the competency and credibility of witnesses” and the “admissibility of a particular piece of evidence.” The Attorney-client privilege, however, is not a rule of competency or credibility. Nor can it be classified as a procedural rule.<sup>10</sup>

\*9 Because the First Restatement does not address this issue and because these jurisdictions do not appear to have otherwise established rules for assessing privilege choice of law matters, the Court believes that the proper conflicts rule and the rule which would be applied by these jurisdictions is section 139 of the Second Restatement.

**e. Section 139 of the Second Restatement—relevant considerations**

The Second Restatement provides that the state with the most significant relationship with the communication is usually the state where the communication took place, *i.e.*, “the state where an oral interchange between persons occurred, where a written statement was received or where an inspection was made of a person or thing.”[Restatement \(Second\) of Conflict of Laws § 139](#), Comment e (1971).

Further, under the Second Restatement, if a communication is privileged in the state to which it bears the most significant relationship, but not privileged in the forum, the law of the state with the most significant relationship to the communication will apply if there is a special reason not to apply the law of the forum. [See Restatement \(Second\) of Conflict of Laws § 139](#). Factors to be considered in assessing whether a special reason exists include:

- (1) the number and nature of the contacts that the state of the forum has with the parties and with the transaction involved, (2) the relative materiality of the evidence that is sought to be excluded, (3) the kind of privilege involved and

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(4) fairness to the parties.

**Restatement (Second) of Conflict of Laws § 139, Comment d (1971).** In addition, where the contacts with the forum are “few and insignificant” then the forum will likely apply the law of the state with the most significant relationship with the communication. *Id.* Finally, “fairness to the parties is another important consideration.” Fairness to the parties requires consideration of the parties’ reasonable expectations of confidentiality which were likely based on the law of the state which has the most significant relationship to the communication. *See Id.*

Considering the above, in the Court’s view, where the communication at issue is between individuals foreign to the forum, whose relationship is centered outside of the forum, and whose communications regard subject matter not centered in the forum, a special reason exists for not applying the law of the forum and the law of the state with the most significant relationship to the communication should be applied. Accordingly, in the instant case, where state privilege law applies, the Court expects that in most if not all instances the law of the state with the most significant relationship to the communication will govern the existence and scope of attorney-client privilege.

#### **f. State Choice of Law Principles**

Courts in thirteen states, the District of Columbia, and Puerto Rico, have adopted section 139 and/or have cited with approval provisions of section 139 when resolving privilege choice of law issues:

**\*10 1. Colorado:** See *People v. Thompson*, 950 P.2d 608, 611 (Colo.App.1997) (concluding that section 139 provided the appropriate framework for analyzing the issue of marital privilege).

**2. Delaware:** See *3Com Corp. v. Diamond II Holdings, Inc.*, 2010 WL 2280734, \*5 (Del. Ch.2010 May 31, 2010) (not reported) (applying section 139 to an attorney-client privilege issue).

**3. District of Columbia:** See *Independent Petrochemical Corp. v. Aetna Cas. and Sur. Co.* 117 F.R.D. 292, 295–296 (D.D.C.1987) (because the District of Columbia typically applies an “interest analysis” approach and relies on the Second Restatement for other choice of law matters it would likely adopt section 139 for privilege matters).

**4. Illinois:** See *Allianz Ins. Co. v. Guidant Corporation*,

869 N.E.2d 1042, 1048–1049 (Ill.App.2007) (section 139 governs issue of attorney-client privilege); *Sterling Finance Management, L.P. v. UBS PaineWebber, Inc.*, 782 N.E.2d 895, 903–904 (Ill.App.2002) (same).

**5. Iowa:** See *State v. Eldrenkamp*, 541 N.W.2d 877 (Iowa 1995) (looking to section 139 for guidance on privilege issue).

**6. Kentucky:** See *Saleba v. Schrand*, 300 S.W.3d 177, 181–183 (Ky.2009) (applying section 139 to privilege issue).

**7. Maine:** See *State v. Lipham*, 910 A.2d 388, 392 n. 3 (Me.2006) (considering section 139 when assessing choice of law issue regarding marital privilege).

**8. Ohio:** See *Woefling v. Great-West Life Assur. Co.*, 285 N.E.2d 61, 221 n. 2 (Ohio App.1972) (finding that Illinois physician-patient privilege controlled and citing to § 139).

**9. Minnesota:** See *State v. Heaney*, 689 N.W.2d 168, 175–177 (Minn.2004) (applying the most significant relationship approach of section 139 to privilege choice of law analysis).

**10. New York:** See *Brandman v. Cross & Brown Co. of Florida, Inc.*, 479 N.Y.S.2d 435, 436–437 (N.Y.Sup.1984) (referencing section 139 and stating that the attorney-client privilege is substantive for purposes of choice of law and New York courts will apply the law of the state with the more significant contacts). See also *Mazella v. Philadelphia Newspapers, Inc.*, 479 F.Supp. 523 (D.C.N.Y.1979) (Neaher, J.) (applying New York choice of law principles; considering section 139 and applying Pennsylvania privilege law because the communication was centered in Pennsylvania).

**11. Pennsylvania:** See *James Talcott, Inc. v. C.I.T. Corp.*, 14 pa. D & C.3d 204, 206 (Pa.Com.Pl.1980) (referencing section 139 and applying the accountant-client privilege law of the state with the most significant relationship to the communication). See also *Samuelson v. Susen*, 576 F.2d 546, 551 (3rd Cir.1978) (Pennsylvania courts have adopted the “interest analysis” approach to conflict questions and therefore would apply the privilege law of the state with the most significant relationship to the privileged communication—particularly when there was no connection between the communication and the forum).

**\*11 12. Puerto Rico:** See *Mitsui & Co. (U.S.A.) Inc.*

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v. Puerto Rico Water Resources Authority, 79 F.R.D. 72 (D.C. Puerto Rico) (Toledo, C.J.) (considering relevant case law and section 139 of the Restatement; concluding that New York law governed the issue of accountant privilege because New York was the state with the most significant relationship to the privileged communication).

**13. Texas:** See *Alez v. State*, 45 S.W.3d 101, 103–106 (Tex.Crim.App.2001) (applying section 139 to privileged communication issue).

**14. Washington:** See *State v. Donahue*, 18 P.3d 608, 611 (Wash.App.Div.2001) (applying section 139 to physician-patient privilege);

**15. Wisconsin:** See *State v. Kennedy*, 396 N.W.2d 765, 769–770 (Wis.App.1986) (relying on section 139 of the Restatement (Second) and concluding that Wisconsin's physician-patient privilege controlled).

As to the jurisdictions that do not appear to have considered privilege issues in the choice of law context, research indicates that twenty-three look to the Second Restatement for guidance when resolving choice of law questions in one or more other areas of law:

**1. Alaska:** See *Savage Arms, Inc. v. Western Auto Supply Co.*, 18 P.3d 49, 53 (Alaska 2001) (generally, Alaska courts look to the Second Restatement in resolving choice of law issues).

**2. Arizona:** See *Bryant v. Silverman*, 703 P.2d 1190, 1191–1192 (generally, Arizona has adopted the Second Restatement for choice of law issues).

**3. Connecticut:** *Jaiguay v. Vasquez*, 948 A.2d 955, 972–973 (Conn.2008) (Connecticut has adopted the significant relationship test of the Second Restatement for personal injury actions noting that “[c]hoice of law must not be rendered a matter of happenstance, in which the respective interests of the parties and the concerned jurisdictions receive only coincidental consideration).

**4. Florida:** See *Bishop v. Florida Specialty Paint Co.*, 389 So.2d 999, 1000–1001 (Fla.1980) (adopting the significant relationship test of Second Restatement for personal injury actions and noting that the court would not apply the law of a state with no connection to the issue). See also *Anas v. Blecker*, 141 F.R.D. 530, 531–533 (M.D.Fla.1992) (Wilson, J.) (concluding that because Florida courts consistently look to the Second Restatement in resolving conflict issues, Florida courts would also apply section 139 to resolve choice of law

issues regarding privilege); *Wolpin v.. Philip Morris Inc.*, 189 F.R.D. 418, 424 (C.D.Cal.1999) (Florida courts would apply section 139 to privilege conflict of law issue) (Paez, J.).

**5. Idaho:** See *Seubert Excavators, Inc. v. Anderson Logging Co.*, 889 P.2d 82, 85 (Idaho 1995) (“Although never adopted in full, this Court has opted in favor of applying the most significant relationship test set forth in the [Second Restatement].”).

**6. Massachusetts:** See *Cosme v. Whitin Mach. Works, Inc.*, 632 N.E.2d 832, 834 (Mass.1994) (in general, choice of law matters are resolved by considering various “choice influencing considerations,” including those provided in the Second Restatement).

**\*12 7. Michigan:** See *Compuware Corp. v. Moody's Investors Services, Inc.*, 222 F.R.D. 124, 130–131 (E.D.Mich.2004) (Feikens, J.) (noting that Michigan has adopted the Second Restatement for contract issues and a two-step interest analysis in tort cases then applying both section 139 and Michigan's two-step interest analysis to resolve disputed privilege issues)

**8. Mississippi:** See *McDaniel v. Ritter*, 556 So.2d 303, 310 (Miss.1989) (Second Restatement applies to torts); *Zurich American Ins. Co. v. Goodwin*, 920 So.2d 427, 433 (Miss.2006) (Second Restatement applicable to contracts). See also *Barnes v. A Confidential Party*, 628 So.2d 283, 288 (Miss.1993) (indicating a preference for application of the law of the state where communication took place).

**9. Missouri:** See *Kennedy v. Dixon*, 439 S.W.2d 173, 184–185 (Mo.1969) (adopting section 145 of the Second Restatement); *Goede v. Aerojet General Corp.*, 143 S.W.3d 14, 25–26 (Mo.App. E.D.2004) (applying the significant relationship test of the Second Restatement); *Byers v. Auto-Owners Ins. Co.*, 119 S.W.3d 659, 663 (Mo.App.S.D.2003) (applying sections 188 and 193 of the Second Restatement).

**10. Montana:** See *Phillips v. General Motors Corp.*, 995 P.2d 1002, 1007–1007 (Mont.2000) (adopting Second Restatement for tort issues); *Tenas v. Progressive Preferred Ins. Co.*, 197 P.3d 990, 994–995 (Mont.2008) (as to contract issues, Montana courts generally follow the Second Restatement).

**11. Nebraska:** See *Erickson v. U-Haul Intern.*, 767 N.W.2d 765, 772–773 (Neb.2009) (Nebraska looks to the Second Restatement for guidance when resolving choice of law issues).

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**12. New Hampshire:** See *Głowski v. Allstate Ins. Co.*, 589 A.2d 593, 595 (N.H.1991) (looking to the Second Restatement when “choice influencing considerations” do not provide enough guidance).

**13. New Jersey:** See *P.V. ex rel. T.V. v. Camp Jaycee*, 962 A.2d 453, 455 (N.J.2008) (applying the most significant relationship standard of the Second Restatement to tort related issue).

**14. Nevada:** *General Motors Corp. v. Dist. Ct.*, 134 P.3d 111, 116 (Nev.2006) (holding that the “Second Restatement’s most significant relationship test governs choice-of-law issues in tort actions unless another, more specific section of the Second Restatement applies to the particular tort”).

**15. Oklahoma:** See *Bernal v. Charter County Mut. Ins. Co.*, 209 P.3d 309, 315 (Okla.2009) (generally lex loci rules are applied in contract actions, however, where motor vehicle insurance contracts are involved, Oklahoma courts will consider whether another state has a more significant interest); *Brickner v. Gooden*, 525 P.2d 632 (Okla.1974) (adopting the most significant relationship test of the Second Restatement in tort actions).

**16. Oregon:** See *Spirit Partners, LP v. Stoel Rivers LLP*, 157 P.3d 1194, 1200 (Or.App.2007) (applying the most significant relationship test of the Second Restatement to tort claims); *M*

**W Zander, U.S. Operations, Inc. v. Scott Co. of California**, 78 P.3d 118, 121 (Or.App.2003) (applying the Second Restatement to contract claim).

**\*13 17. Rhode Island:** See *Najarian v. National Amusements, Inc.*, 768 A.2d 1253, 1255 (R.I.2001) (applying Rhode Island’s “interest-weighing” approach in deciding choice of law issue but also looking to Second Restatement for guidance).

**18. South Dakota:** See *Burhenn v. Dennis Supply Co.*, 685 N.W.2d 778, 784 (S.D.2004) (South Dakota applies the significant relationship test of the Second Restatement to resolve choice of law concerns);

**19. Tennessee:** See *Hataway v. McKinley*, 830 S.W.2d 53, 59 (Tenn.1992) (Tennessee will follow the “most significant relationship” approach of the Second Restatement for conflicts questions in tort cases).

**20. Utah:** See *Waddoups v. Amalgamated Sugar Co.*, 54 P.3d 1054, 1059 (Utah 2002) (Utah applies the Second Restatement when resolving choice of law

issues). See also *Hercules, Inc. v. Martin Marietta Corp.*, 143 F.R.D. 266, 268–269 (D.Utah 1992) (predicting that Utah would look to section 139 for assessing privilege choice of law issues because Utah has applied the significant contacts analysis of the Second Restatement in other areas of law).

**21. Vermont:** See *McKinnon v. F.H. Morgan Co., nc.*, 750 A.2d 1026, 1028–128 (Vt.2000) (in Vermont, most significant relationship test of Second Restatement is applied to tort claims); *Pioneer Credit Corp. v. Carden*, 245 A.2d 891, 894 (Vt.1968) (applying Second Restatement to resolve choice of law issue pertaining to contract).

**22. West Virginia:** See *Lee v. Saliga*, 373 S.E.2d 345, 351–352 (W.Va.1988) (discussing West Virginia’s history of applying the Second Restatement to contract matters).

**23. Wyoming:** See *R & G Elec., Inc. v. Devon Energy Corp.*, 53 Fed. Appx. 857, 859 (10th Cir.2002) (Wyoming follows the Second Restatement approach in resolving choice of law questions).

Research indicates that another six jurisdictions apply an interest analysis that reflects principles found in the Second Restatement when addressing other choice of law matters.

**1. Arkansas:** See *Crisler v. Unum Insurance Co. of America*, 233 S.W.3d 658, 660 (Ark.2006) (in contract dispute applying the law of the state with the most significant relationship); *Ganey v. Kawasaki Motors Corp., U.S.A.*, 234 S.W.3d 838, 846–847 (Ark.2006) (choice of law matters in tort are resolved by considering both the doctrine of lex loci delecti and the five “choice-influencing considerations” promulgated by Professor Robert A. Leflar; in rendering its decision the court also considered which state had a more significant relationship to the issue in question).

**2. California:** See *Wolpin v. Philip Morris Inc.*, 189 F.R.D. 418, 423–424 (D.D.Cal.1999) (Paez, J.) (applying California’s governmental interest analysis to determine which state had the most significant connection with a confidential study; considering factors such as residence of study participants and involvement of California Department of Health Services in collecting data).

**\*14 3. Hawaii:** See *Peters v. Peters*, 634 P.2d 586, 593 (Hawaii 1981) (“The preferred analysis, in our

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opinion, would be an assessment of the interests and policy factors involved with a purpose of arriving at a desirable result in each situation.”).

**4. Indiana:** See *Hubbard MFG. Co., Inc. v. Greeson*, 515 N.E.2d 1071 (Ind.1987) (in torts Indiana follows a modified version of the “lex loci delicti” rule; if the place of the tort is an insignificant contact the court may consider other factors including the place where the relevant relationship is centered); *Hartford Acc. Indem. Co. v. Dana Corp.*, 690 N.E.2d 285 (Ind.Ct.App.1997) (in contract actions Indiana applies the law of the state with the most significant contacts).

**5. Louisiana:** See Article 3515 of the Louisiana Civil Code codifies a choice of law analysis that considers each states relationship to the parties and the dispute, including consideration of the “policies of upholding the justified expectations of parties”).

**6. North Dakota:** See *Johnson v. Johnson*, 617 N.W.2d 97, 123 (N.D.2000) (applying a significant contacts approach coupled with “choice influencing considerations” to contracts choice of law issues); *Nodak Mut. Ins. Co. v. Wamsley*, 687 N.W.2d 226, 231 (N.D.2004) (applying Second Restatement to choice of law issues in tort).

Finally, the Court’s research reveals eight states who continue to follow the traditional choice of law principles largely reflected in the original Restatement (First) of Conflict of Laws (1934) (“First Restatement”).

**1. Alabama:** See *Lifestar Response of Alabama, Inc. v. Admiral Ins. Co.*, 17 So.3d 200, 213 (Ala.2009) (Alabama generally follows conflict of law principles of *lex loci contractus* and *lex loci delicti* ).

**2. Georgia:** See *Convergys Corp. v. Keener*, 582 S.E.2d 84 (Ga.2003) (declining to adopt the Second Restatement and continuing to apply a more traditional approach).

**3. Kansas:** See *Aselco, Inc. v. Hartford Ins. Group*, 21 P.3d 1011, rev. denied 272 Kan.— (2001) (applying traditional choice of law analysis).

**4. Maryland:** See *Hauch v. Connor*, 453 A.2d 1207, 1209 (Md.1983) (declining to adopt the Second Restatement for tort choice of law matters); *American Motorists Ins. Co. v. Artra Group, Inc.*, 659 A.2d 1295, 1301 (Md.1995) (Maryland has not adopted the significant relationship test of the Second Restatement but has cited with approval

other provisions of the Second Restatement). The United States District Court for the District of Maryland has, on two occasions, concluded that, as to matters of privilege, Maryland courts would apply section 139 of the Second Restatement). See *Saint Annes Development Co., LLC v. Trabich*, 2009 WL 324054, \*2 (D.Md. Feb. 9, 2009) (Bredar, Mag.); *Hill v. Huddleston*, 263 F.Supp. 108 (D.Md.1967) (Thomsen, C.J.).

**5. New Mexico:** See *United Wholesale Liquor Co. v. Brown-Forman Distillers Corp.*, 775 P.2d 233, 235 (1987) (noting “New Mexico adheres to a traditional conflicts of law analysis contained in Restatement (First) of Conflicts of Law (1934)”). But c.f. *Ideal v. Burlington Resources Oil & Gas Co. LP*, 233 P.3d 362, 369 (N.M.2010) (noting that the New Mexico Supreme Court recently rejected the First Restatement approach as too rigid and inflexible in multi-state contract class actions; adopting instead the “more appropriate” Second Restatement approach).

\*15 **6. North Carolina:** See *Gbye v. Gbye*, 503 S.E.2d 434, 435–436 (N.C.1998) (North Carolina adheres to a traditional choice of law approach in tort actions); *Computer Sales Intern., Inc. v. Forsyth Memorial Hosp., Inc.*, 436 S.E.2d 263, 265 (N.C.App.1993) (law of the place where the contract is made governs its interpretation)

**7. South Carolina:** See *Menezes v. WL Ross Co. LLC*, — S.E.2d —, 2011 WL 1118841, 3 (S.C.App.2011 March 23, 2011)

**8. Virginia:** See *Jones v. R.S. Jones & Assocs.*, 431 S.E.2d 33, 34 (Va.1993) (“Virginia applies the doctrine of lex loci delicti, meaning the law of the place of the wrong governs all matters related to the basis of the right of action.”); *Buchanan v. Doe*, 431 S.E.2d 289, 291 (Va.1993) (under Virginia “conflict of law rules: (1) the law of the place of the wrong determines the substantive issues of tort liability, and (2) generally, the law of the place where an insurance contract is written and delivered controls issues as to its coverage.”) (internal citations omitted).

#### **IV. CONCLUSION AND SUMMARY OF FINDINGS**

Having considered the parties’ arguments, the relevant authority, and for the reasons stated herein, the Court ORDERS as follows:

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1. Federal law will govern whether documents are protected under the work-product doctrine.
2. The confidentiality of communications that are relevant to an element of a federal defense will be governed by federal attorney-client privilege law.
3. The confidentiality of communications relevant to an element of a claim or defense for which state law supplies the rule of decision will be governed by state privilege law.
4. The confidentiality of communications relevant to both federal and state elements will be governed by the law favoring reception of the evidence.
5. Choice of law considerations for matters governed by state privilege law
  - a. Section 139 of the Second Restatement will govern the choice of law decision for matters governed by state privilege law.
  - b. As the Court interprets section 139 of the Second Restatement, vindication of the parties' reasonable expectations of confidentiality is a "special circumstance" warranting application of the law of the state with the most significant relationship to the communication. Thus, where the communications at issue are between individuals foreign to the forum, whose relationship is centered outside of the forum, and whose communications regard subject matter not

**Footnotes**

<sup>1</sup> Professor Paul Rice, the attorney-client privilege expert appointed in the *Vioxx* litigation, addresses this concept in his treatise on attorney-client privilege law in the United States. See 2 Paul R. Rice & John B. Corr, *Attorney-Client Privilege in the U.S.* § 12:17 (2010) (noting that Rule 501's directive to apply state privilege law "intersects with the *Erie* doctrine" and requires the application of state choice of law rules in determining which state's privilege law controls).

<sup>2</sup> In addition, the agreed order entered in this MDL that allows for direct filing expressly states that the parties' agreement to permit direct filing would not impact the choice of law that otherwise would apply to the direct filed actions (MDL 2100 Docs 669 (original direct filing order), 1137(amended direct filing order), & 1462 (second amended direct filing order)).

<sup>3</sup> The Advisory Committee Notes that accompany Rule 501 expressly state ("[in] Federal question civil cases, federally evolved rules on privilege should apply since it is Federal policy which is being enforced. *It is also intended that the Federal law of privileges should be applied with respect to pendent State law claims when they arise in a Federal question case*"). Fed.R.Evid. 501, Advisory Committee Notes, S.Rep. No. 93-1277 (emphasis added). A number of courts have relied on this statement in concluding that federal question cases involving pendent state claims are governed by federal privilege law. This statement, however, is contained in the senate report that accompanies Rule 501 and was made in reference to the senate's version of Rule 501. The Conference Report is silent on the issue and, as a result, congressional intent is unclear. Accordingly, as to the issue of what law governs pendent state law claims in federal question cases, the Court relies on Seventh Circuit authority and not the commentary contained in senate report No. 93-1277.

<sup>4</sup> This portion of the Conference Report appears to reject the position taken by the Senate Judiciary Committee regarding what constitutes an "element" of a claim or defense. See 23 Charles Alan Wright & Kenneth Graham, Jr., *Federal Practice And Procedure: Evidence* § 5434 (pursuant to the Conference Report "it makes no difference whether the supposedly privileged matter is direct or circumstantial evidence of a state claim; if it is in a line of proof that culminates in an element of a state claim or

centered in the forum, the forum's privilege law is not controlling.

6. The parties are directed to meet and confer as specified in section V below.

**V. MEET AND CONFER PROCESS**

In light of the Court's decision resolving the choice of law considerations, the Court directs the parties to meet and confer, for a period not to exceed seven days, with respect to the confidentiality of the 330 disputed documents. The parties should attempt to independently reach an agreement regarding the 330 disputed documents by applying the principles discussed in this order. If the parties are unable to resolve this matter independently, the parties shall submit briefing relevant to any remaining disputes over the 330 disputed documents. The briefing shall include: (1) a list of documents as to which disputes remain over claims of attorney-client privilege and work-product; (2) position statements addressing the disputes which remain over claims of attorney-client privilege and work-product; and (3) defendants' privilege logs.

**\*16 SO ORDERED.**

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defense, then state rules of privilege apply").

5 This is not the first time this issue has arisen in this MDL. The Court has explained, in previous MDL decisions, that transfer cases are not governed by Illinois choice of law principles. *See e.g., In re Yasmin and Yaz (Drospirenone) Marketing, Sales Practices and Relevant Products Liability Litigation*, 2010 WL 3937414, \*4 (S.D.Ill., Oct. 4, 2010) (Herndon, C.J.); *In re Yasmin and Yaz (DROSPIRENONE) Marketing, Sales Practices and Products Liability Litigation*, 2010 WL 1963202, \*2 (S.D.Ill. May 14, 2010) (Herndon, C.J.).

6 MDL 2100 Docs 669 (original direct filing order), 1137(amended direct filing order), & 1462 (second amended direct filing order).

7 Jurisdiction in the *Vioxx* litigation was diversity based and the coordinated proceedings involved cases that originated in jurisdictions across the country. *Vioxx*, 501 F.Supp.2d at 789–790.

8 The House Report also states “where an element of a claim or defense is not grounded upon a federal question, there is no federal interest strong enough to justify departure from State policy.”*Fed.R.Evid. 501* Advisory Committee Notes, H.R. No. 93–650.

9 The Court provides a detailed review of the relevant case law from these jurisdictions in section III(B)(2)(f) below.

10 The Court notes that some courts have concluded that the issue of attorney-client privilege is procedural in nature and have therefore applied the law of the forum. *See e.g., Union Planters Nat'l Bank v. ABC Records, Inc.*, 82 F.R.D. 472 (W.D.Tenn.1979). The Court does not rely on these decisions because *Federal Rule of Evidence 501* rejects the notion that the attorney-client privilege issue is procedural. *Fed.R.Evid. 501* Advisory Committee Notes, H.R. 93–650. As was well said by the Second Circuit Court of Appeals in *Republic Gear Co. v. Borg-Warner Corp.*, 381 F.2d 551, 555 n. 2 (2d Cir.1967):

[A] rule of privilege is unlike the ordinary rules of practice which refer to the processes of litigation, in that it affects private conduct before the litigation arises. Rules of privilege are not mere housekeeping rules which are rationally capable of classification as either substantive or procedural for purposes of applying the doctrine of [Erie ]. Such rules ‘affect people’s conduct at the stage of primary private activity and should therefore be classified as substantive or quasi-substantive.

(internal citations omitted). The Court also notes that if it were to treat any state’s attorney-client privilege as merely procedural that state’s privilege law could not be applied because federal courts sitting in diversity are required to adhere to federal rules of procedure.

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